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(21) International Application Number: PCT/US97/02183 (22) International Filing Date: 11 February 1997 (11.02.97) (30) Priority Data: 08/600,058 12 February 1996 (12.02.96) US (71) Applicant: HEALTHPOINT, LTD. [US/US]; 307 E. Josephine Street, San Antonio, TX 78215-1128 (US). (72) Inventors: MINER, Norman, A.; Healthpoint, Ltd., 307 E. Josephine Street, San Antonio, TX 78215-1128 (US). WOLLER, William, H.; Healthpoint, Ltd., 307 E. Josephine Street, San Antonio, TX 78215-1128 (US). ANDERSON, Edward, L.; Healthpoint, Ltd., 307 E. Josephine Street, San Antonio, TX 78215-1128 (US). HOBSON, David, W.; Healthpoint, Ltd., 307 E. Josephine Street, San Antonio, TX 78215-1128 (US). (74) Agent: SEASE, Edmund, J.; Zarley, McKee, Thome, Voorhees & Sease, Suite 3200, 801 Grand Avenue, Des Moines, IA 50309-2721 (US).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: QUICK ACTING CHEMICAL STERILANT (57) Abstract A low odor, aqueous, quick acting room temperature disinfectant slution primarily useful for medical instruments to disinfect within a half hour or less. The composition comprises a reacting or synergistic combination of hydrogen peroxide and from about 1 % to 30 % by weight of a water soluble organic acid or salt form thereof with the acid preferably being selected from the group consisting of malonic and succinic acids.		

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TITLE: QUICK ACTING CHEMICAL STERILANT

BACKGROUND OF THE INVENTION

Medical, dental and other instruments are often made of high quality stainless steel that can be cleaned and sterilized between uses for different patients by high temperature steam under pressure. This sterilization procedure is quick, reliable, odorless, non-toxic and inexpensive. In contrast to this situation, more and more instruments are now made of heat-sensitive plastic, rubber, glass lenses and electronic components. These flexible, flexible-lensed, and rigid-lensed instruments allow relatively non-invasive diagnostic and treatment procedures within the body. The non-invasive procedures allowed by these heat-sensitive instruments are responsible for great advances in medical practice. During use, these instruments can be contaminated with deadly pathogens such as the Human Immunodeficiency Virus (HIV), hepatitis viruses, and antibiotic drug-resistant tuberculosis and other bacteria. For these reasons, it is imperative that these heat-sensitive instruments be sterilized of all microbes prior to each use. The chemical germicides available for sterilization of heat-sensitive instruments have in the past had many problems that made their use difficult.

The antimicrobial properties of hydrogen peroxide have been known for many years. However, 6% hydrogen peroxide requires a minimum of 6 hours at room temperature to pass the standard Association of Official Analytical Chemists (AOAC) Sporocidal Test. This is the test that defines "sterilant" for liquid chemical germicides in the United States. The antimicrobial properties of peracetic acid are also well known. Peracetic acid has a very sharp pungent odor, and is known as a tumor-promoting agent when tested on mouse skin. For these reasons, the use of peracetic acid as a chemical sterilant is limited to low concentrations used with enclosed systems.

Antimicrobial synergism between hydrogen peroxide and peracetic acid is a well established fact. Such compositions are prepared by mixing hydrogen peroxide and acetic acid to give equilibrated solutions of hydrogen peroxide, acetic acid, and peracetic acid. There is a great deal of scientific and patent literature regarding hydrogen peroxide-peracetic acid solutions for sterilization. By way of example only, Minntech Corporation of Minneapolis, Minnesota, has a kit or sterilization console for disinfecting with hydrogen peroxide-peracetic acid solutions (U.S. Patent 5,400,818). However, this combination is limited by the same problems of pungent odor and potential toxicity as peracetic acid alone. This often means that such formulations are used at such dilute concentrations that rapid sporicidal activity is lost, or the solutions are limited to enclosed systems that contain the pungent fumes.

Steris Corporation of Mentor, Ohio, markets a Steris System 1 product. This uses a low concentration of peracetic acid (about 0.2%) contained within a machine, and is heated to 122°F to achieve rapid sterilization. The relatively low peracetic acid concentration, coupled with the high temperature, breaks down the peracetic acid, limiting it to one single use cycle. The heated, enclosed, single-use machine system is expensive and less than desirable.

Another chemical sterilant is 2% alkaline glutaraldehyde. Glutaraldehyde requires about 10 hours at 25°C to pass the AOAC Sporicidal Test. Because of this long exposure time, the use of glutaraldehyde is usually compromised to accept disinfection from a shorter exposure time rather than the safer condition of sterilization. Furthermore, glutaraldehyde has an odor that irritates eye, nose, and throat mucous membranes. Repeated exposure to glutaraldehyde causes headaches and allergic reactions for some people. For these reasons, glutaraldehyde is a less than desirable chemical germicide.

Many chemicals that contain chlorine are rapidly sporicidal and capable of sterilization. Examples are

bleach, the active agent of which is HOCl , HClO_2 , ClO_2 , and HCl . However, while these chemicals are rapidly sporicidal, they are too corrosive to metals and elastomers to find any practical use in sterilization of medical, dental and other instruments.

It can therefore be seen that there is a continuing need for an effective, practical, safe, affordable sterilant for heat-sensitive instruments, as well as for all applications that are beyond the scope of steam sterilization. This invention has as its primary objective the fulfillment of this need.

SUMMARY OF THE INVENTION

This invention relates to a rapid acting room temperature sterilant. It is a low odor, aqueous disinfecting solution having a pH within the range of 2-6. It comprises in combination a solution of from about 1% to about 30% by weight of peroxide capable of releasing hydroxyl free radicals, and from about 1% to about 30% by weight of a water soluble organic acid or salt form of a C_3 or higher mono, or a di-, tri-, or poly carboxylic organic acid, with the organic acid preferably selected from the group consisting of malonic acid and succinic acid, or combinations thereof. It is believed there may be a reaction between the peroxide and carboxylic acids that produce a third chemical or condition that causes rapid kill of bacterial spores and other microbes at ambient temperatures (18°C - 24°C) in short times (i.e. within 30 minutes). The carboxylic acids that can be used with peroxides can be selected from a large group to be relatively odor-free, non-toxic, soluble and inexpensive.

DETAILED DESCRIPTION OF THE INVENTION

The sterilizing and disinfecting solutions of this invention have a variety of uses. The solutions have excellent sterilization and disinfecting properties and can be used to sterilize sophisticated medical instruments such

as endoscopes without causing damage to sensitive parts of such instruments.

The fact that this process can be used with endoscopic instruments is significant since relatively non-invasive
5 endoscopic procedures have revolutionized the way that surgery is performed. As earlier mentioned, few rigid or flexible endoscopes can be sterilized by the quick and sure method of steam sterilization because the plastic, rubber and precisely-positioned glass lenses of
10 endoscopes make them incompatible with the heat of a steam sterilizer. Instead they must be sterilized using lower temperatures and typically slower processes. They also must use a sterilizing solution that is non-corrosive.

Endoscopes are but one example of the type of instrument
15 that can be effectively sterilized with the present compositions. Conventional surgical instruments of all types, microsurgery instrument sets, anesthesia equipment, etc. can also be treated. Generally, the composition disclosed herein can be used for sterilization of any
20 products that enter sterile tissue or the vascular system or have tissue contact during any surgeries. Necessarily, if the solution is effective for these critical medical instruments, it also can be used for intermediate level and low level instruments and surfaces. Because the formulation
25 is relatively odorless and non-toxic, one can sterilize surfaces that formerly were only disinfected or sanitized, or one can dilute the formula for disinfection rather than sterilization. The composition may also be used as an antiseptic to kill germs on skin. It is therefore versatile
30 in use.

It had previously been thought that effectiveness of hydrogen peroxide and peracetic acid combinations to pass standard "AOAC" sterilization tests was due to the
substantial enhancement of formation of free hydroxy radicals
35 from the "per" acid in combination with hydrogen peroxide. Accordingly, it was thought necessary to add peroxyacetic acid directly to germicidal formulations. Quite

surprisingly, the inventors found that it was not necessary to add the toxic and malodorous peracetic acid to sterilizing formulations. Instead, certain lower carboxylic acids selected for their solubility, lack of odor, and non-toxic nature can be used in combination with hydrogen peroxide to achieve sterility at ambient temperatures and short exposure times. In particular, the carboxylic acid could be a C₃ or higher mono or a di- or poly carboxylic acid of up to C₁₂ chain length and can be saturated or unsaturated. As a result of this composition, complex and expensive equipment needed to contain toxic chemicals can be eliminated, and since the acids employed are weak organic acids, corrosion of materials is significantly reduced. As a result, providing the levels herein described are used, there is a reacting or synergistic relationship between the defined water soluble organic acid and the peroxide such that even at lower temperatures non-corrosive sterilization is achieved. Moreover, the chemicals are generally inexpensive and odor-free, and are therefore economically and simply packaged. Of course, less odor and less toxic mean that higher concentration can be used with accompanying faster rate of sterilization. At higher exposure temperatures of 30, 40, or 50°C, for example, the exposure time needed to achieve sterilization is even faster than at ambient temperatures.

The first component of the composition is from about 1% to about 30% by weight of a peroxide. Preferably, the amount of peroxide is from about 1% by weight to about 12% by weight of the disinfecting solution, and most preferably from about 6% by weight to about 10% by weight of the disinfecting solution. The preferable concentration of peroxide may be varied depending on the application from lower concentration for an antiseptic to higher concentrations for a low-temperature, rapid-acting sterilant. The peroxide of choice is, of course, the most commonly available peroxide, hydrogen peroxide. However, the invention is not limited to hydrogen peroxide, and other peroxy compounds may be employed. These include, for example, perborates, saturated and unsaturated

peralkanoic acids such as peracetic acid, performic acid, perpropionic acid, etc. The critical factor is that it be a water soluble peroxide compound that is compatible with the weak carboxylic acid component.

5 The weak carboxylic acid component of the present invention is preferably a di- acid of lower C_{12} or less carbon length carboxylic acid preferably selected from the group consisting of malonic acid and succinic acid. Also, examples of acids in this class would be malic, oxalic and
10 tartaric acids. These acids, when in the proper concentrations, are low odor, reasonably soluble and non-corrosive. The amount of the carboxylic acid component generally would be within the range of from about 1.0% by weight to 30% by weight of sterilizing or disinfecting
15 solution, preferably from about 1% by weight to about 12% by weight of the solution, and most preferably from about 3% by weight to about 6% by weight of the solution composition. As with the peroxide, the preferred concentration of carboxylic acids is related to the intended end use.

20 Generally speaking, and as a guideline, the peroxide component should have a concentration of within the range of 0.2M to about 10M, preferably within the range of 0.2M to 4.0M. The organic acid component should have a concentration within the range of 0.05M to 4.0M, and preferably of 0.05M to
25 2.0M.

While acetic acid is unacceptable by itself because of its normal pungent odor, it is possible that some acetic acid, in combination with other of the acids described here, can be successfully used. Thus the key to the present
30 invention is the presence of the herein-described combination or perhaps the reaction product thereof.

Generally, the amount of peroxide component and the amount of carboxylic acid component are balanced such that the pH will be within the range of about 2.0 to 6.0,
35 preferably about 3.0 to 5.0.

While a suitable sterilizing and disinfecting solution can be achieved with these two components only, as is

understood by those skilled in the art, other ingredients may be added. In fact, the sterilizing and disinfecting capabilities can be enhanced by adding a small amount of detergent such as nonionic or anionic detergent. The amount
5 of detergent can be within the range of from about 0.05% by weight to about 1.0% by weight, preferably from about 0.1% by weight to about 0.5% by weight. The amount of detergent should be enough to enhance the sterilization and disinfection, but less than the amount which would provide
10 substantial sudsing.

Suitable synthetic detergents are well known to those of ordinary skill in the art, but generally these surface active agents can be selected from the group consisting of anionic and nonionic surfactants. Non-ionic, ether-linked
15 surfactants such as Laureth®4 or Laureth®23 are preferred.

Alkyl sulfate surfactants are a type of anionic surfactant of importance for use herein. Alkyl sulfates have the general formula ROSO_3M wherein R preferably is a C_{10} - C_{24} hydrocarbyl, preferably an alkyl or hydroxyalkyl having a
20 C_{10} - C_{20} alkyl component, more preferably a C_{12} - C_{18} alkyl or hydroxyalkyl, and M is H or a cation, e.g., an alkali metal cation (e.g., sodium, potassium, lithium), substituted or unsubstituted ammonium cations such as methyl-, dimethyl-, and trimethyl ammonium and quaternary ammonium cations, e.g.,
25 tetramethylammonium and dimethyl piperdinium, and cations derived from alkanolamines such as ethanolamine, diethanolamine, triethanolamine, and mixtures thereof, and the like. Typically, alkyl chains of C_{12} -16 are preferred for lower wash temperatures (e.g., below about 50°C) and C_{16} -
30 18 alkyl chains are preferred for higher wash temperatures (e.g., above about 50°C).

Alkyl alkoxyated sulfate surfactants are another category of useful anionic surfactant. These surfactants are water soluble salts or acids typically of the formula
35 $\text{RO(A)}_m\text{SO}_3\text{M}$ wherein R is an unsubstituted C_{10} - C_{24} alkyl or hydroxyalkyl group having a C_{10} - C_{24} alkyl component, preferably a C_{12} - C_{20} alkyl or hydroxyalkyl, more preferably

C₁₂-C₁₈ alkyl or hydroxyalkyl, A is an ethoxy or propoxy unit, m is greater than zero, typically between about 0.5 and about 6, more preferably between about 0.5 and about 3, and M is H or a cation which can be, for example, a metal cation (e.g., sodium, potassium, lithium, calcium, magnesium, etc.), ammonium or substituted-ammonium cation. Alkyl ethoxylated sulfates as well as alkyl propoxylated sulfates are contemplated herein. Specific examples of substituted ammonium cations include methyl-, dimethyl-, trimethyl-ammonium and quaternary ammonium cations, such as tetramethyl-ammonium, dimethyl piperidinium and cations derived from alkanolamines, e.g., monoethanolamine, diethanolamine, and triethanolamine, and mixtures thereof. Exemplary surfactants are C₁₂-C₁₈ alkyl polyethoxylate (1.0) sulfate, C₁₂-C₁₈ alkyl polyethoxylate (2.25) sulfate, C₁₂-C₁₈ alkyl polyethoxylate (3.0) sulfate, and C₁₂-C₁₈ alkyl polyethoxylate (4.0) sulfate wherein M is conveniently selected from sodium and potassium.

Other anionic surfactants useful for deterative purposes can also be included in the compositions hereof. These can include salts (including, for example, sodium, potassium, ammonium, and substituted ammonium salts such as mono-, di- and triethanolamine salts) of soap, C₉-C₂₀ linear alkylbenzenesulphonates, C₈-C₂₂ primary or secondary alkanesulphonates, C₈-C₂₄ olefinsulphonates, sulphonated polycarboxylic acids, alkyl glycerol sulfonates, fatty acyl glycerol sulfonates, fatty oleyl glycerol sulfates, alkyl phenol ethylene oxide ether sulfates, paraffin sulfonates, alkyl phosphates, isothionates such as the acyl isothionates, N-acyl taurates, fatty acid amides of methyl tauride, alkyl succinamates and sulfosuccinates, monoesters of sulfosuccinate (especially saturated and unsaturated C₁₂-C₁₈ monoesters) diesters of sulfosuccinate (especially saturated and unsaturated C₆-C₁₄ diesters), N-acyl sarcosinates, sulfates of alkylpolysaccharides such as the sulfates of alkylpolyglucoside, branched primary alkyl sulfates, alkyl polyethoxy carboxylates such as those of the formula

RO(CH₂CH₂O)_kCH₂COO-M⁺ wherein R is a C₈-C₂₂ alkyl, k is an integer from 0 to 10, and M is a soluble salt-forming cation, and fatty acids esterified with isethionic acid and neutralized with sodium hydroxide. Further examples are given in *Surface Active Agents and Detergents* (Vol. I and II by Schwartz, Perry and Berch).

Suitable nonionic detergent surfactants are generally disclosed in U.S. Pat. 3,929,678, Laughlin et al., issued Dec. 30, 1975, at column 13, line 14 through column 16, line 6, incorporated herein by reference. Exemplary, non-limiting classes of useful nonionic surfactants are listed below.

The polyethylene, polypropylene, and polybutylene oxide condensates of alkyl phenols. In general, the polyethylene oxide condensates are preferred. These compounds include the condensation products of alkyl phenols having an alkyl group containing from about 6 to about 12 carbon atoms in either a straight chain or branched chain configuration with the alkaline oxide. These compounds are commonly referred to as alkyl phenol alkoxylates, (e.g., alkyl phenol ethoxylates).

The condensation products of aliphatic alcohols with from about 1 to about 25 moles of ethylene oxide. The alkyl chain of the aliphatic alcohol can either be straight or branches, primary or secondary, and generally contains from about 8 to about 22 carbon atoms. Particularly preferred are the condensation products of alcohols having an alkyl group containing from about 10 to about 20 carbon atoms with from about 2 to about 18 moles of ethylene oxide per mole of alcohol.

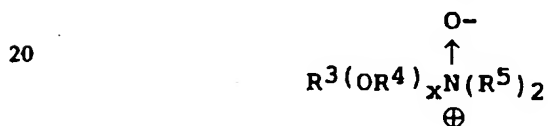
The condensation products of ethylene oxide with a hydrophobic base formed by the condensation of propylene oxide with propylene glycol. Examples of compounds of this type include certain of the commercially-available PluronicTM surfactants, marketed by BASF.

The condensation products of ethylene oxide with the product resulting from the reaction of propylene oxide and ethylenediamine. Examples of this type of nonionic

surfactant include certain of the commercially available Tetronic™ compounds, marketed by BASF.

Semi-polar nonionic surfactants are a special category of nonionic surfactants which include water-soluble amine
 5 oxides containing one alkyl moiety of from about 10 to about 18 carbon atoms and 2 moieties selected from the group consisting of alkyl groups and hydroxyalkyl groups containing from about 1 to about 3 carbon atoms; water-soluble phosphine
 10 oxides containing one alkyl moiety of from about 10 to about 18 carbon atoms and 2 moieties selected from the group consisting of alkyl groups and hydroxyalkyl groups containing from about 1 to about 3 carbon atoms; and water-soluble
 15 sulfoxides containing one alkyl moiety of from about 10 to about 18 carbon atoms and a moiety selected from the group consisting of alkyl and hydroxyalkyl moieties of from about 1 to about 3 carbon atoms.

Semi-polar nonionic detergent surfactants include the amine oxide surfactants having the formula



wherein R³ is an alkyl, hydroxyalkyl, or alkyl phenyl group or mixtures thereof containing from about 8 to about 22
 25 carbon atoms; R⁴ is an alkylene or hydroxyalkylene group containing from about 2 to about 3 carbon atoms or mixtures thereof; x is from 0 to about 3; and each R⁵ is an alkyl or hydroxyalkyl group containing from about 1 to about 3 carbon atoms or a polyethylene oxide group containing from about 1
 30 to about 3 ethylene oxide groups. The R⁵ groups can be attached to each other, e.g., through an oxygen or nitrogen atom, to form a ring structure.

These amine oxide surfactants in particular include C₁₀-C₁₈ alkyl dimethyl amine oxides and C₈-C₁₂ alkoxy ethyl
 35 dihydroxy ethyl amine oxides.

Alkylpolysaccharides disclosed in U.S. Pat. 4,565,647, Llenado, issued Jan. 21, 1986, having a hydrophobic group containing from about 6 to about 30 carbon atoms, preferably

from about 10 to about 16 carbon atoms and a polysaccharide, e.g., a polyglycoside, hydrophilic group containing from about 1.3 to about 10, preferably from about 1.3 to about 3, most preferably from about 1.3 to about 2.7 saccharide units.

5 Any reducing saccharide containing 5 or 6 carbon atoms can be used, e.g., glucose, galactose and galactosyl moieties can be substituted for the glucosyl moieties. (Optionally the hydrophobic group is attached at the 2-, 3-, 4-, etc. positions, thus giving a glucose or galactose as opposed to a glucoside or galactoside). The intersaccharide bonds can be,

10 e.g., between the one position of the additional saccharide units and the 2-, 3-, 4-, and/or 6- positions on the preceding saccharide units.

Fatty acid amide surfactants having the formula:



wherein R⁶ is an alkyl group containing from about 7 to about 21 (preferably from about 9 to about 17) carbon atoms and

20 each R⁷ is selected from the group consisting of hydrogen, C₁-C₄ alkyl, C₁-C₄ hydroxyalkyl, and —(C₂H₄O)_xH where x varies from about 1 to about 3.

In addition to the above, if desired, corrosion inhibitors at very minor levels can be used, i.e. at levels

25 of .01% to .1% on a weight basis. Suitable corrosion inhibitors can include those available and known, for example, complex fatty amine salts such as n,n'dibutylthiourea, etc.

Nonionic ether linked surfactants are preferred such as

30 Laureth®23 or Laureth®4.

In addition to all of the above, as is well understood by those skilled in the art, other minors can be employed to make the basic composition more pharmaceutically elegant. For example, odorants can be added at very minor levels as

35 can dyes, diluents such as alcohol, buffers, etc. With the exception of diluents such as alcohols which are used at

higher levels, the levels of these minors are generally not more than .001% to .01% by weight.

The composition can be used as a sterilant for medical and dental equipment, implanted medical and dental devices and appliances, can be used as a disinfectant for inanimate surfaces, can be used as an antiseptic for skin disinfection, such as for patient preoperative skin disinfection or health personnel, a hand wash, may be used as a disinfectant for contact lenses, an oral disinfectant or antiseptic, and can be used generally for conventional, intermediate and low level disinfection, and as a sterilant in industrial applications.

Packaging of the composition is not complex. It may be prepackaged in dry form if desired with instructions for mixing solutions on the spot, or it may be prepackaged in solution form, ideally in two packages (one the peroxide and one the organic acid component) to be mixed at point of use. This enhances freshness and accuracy of compliance with directions.

The following examples are offered to illustrate, but not limit, the process of the present invention and to demonstrate the surprising result that satisfactory results in comparison with acetic acid can be achieved with weaker longer chain acids such as succinic acid.

Historically, the Environmental Protection Agency regulates germicides in the United States, and the test for a sterilizing claim (a sterilant) by a liquid germicide is the Association of Official Analytical Chemists (AOAC) Sporidical Activity of Disinfectants Test 966.04. This test exposes spores dried onto carrier surfaces to the germicide. To make a label claim as a sterilant, a germicide must produce 720 sterile cylinders of 720 total cylinders within a specified exposure time and temperature range. A legal definition of sterilant in the United States is one that can pass this test. In the following tests peroxide composition alone was compared with an acetic acid composition alone and with sodium acetate composition with regard to ability to

sterilize carriers labeled with spores according to the methods of the AOAC Sporocidal Test.

TABLE I

	Number of Positive (+) Cylinders per Total Number Tested. 30 Min. Exposure at 20±1°C.			Percent Sterile Cylinders
Formulation				
10	6% H ₂ O ₂ . pH 4.7	20/20		Zero
	6% H ₂ O ₂ + 0.5% Acetic Acid pH 2.7	2/20		90%
15	0.5% Acetic Acid pH 2.8	20/20		Zero
	6% H ₂ O ₂ + 0.5% Sodium Acetate pH 6.7	20/20		Zero
20	0.5% Sodium Acetate pH 7.7	20/20		Zero

This test was repeated with some modifications in an attempt to sterilize 100% of the *C.sporogenes*-labeled cylinders. The results were as follows.

TABLE II

	Formulation	Time in Min. at 20±1°C	Number of Positive Cylinders (+) Per Total Number Tested	Percent Sterile Cylinders
35	8% H ₂ O ₂ pH 4.5	30	20/20	Zero
	8% H ₂ O ₂ + 2% Acetic Acid pH 2.4	10 20 30	9/20 0/20 0/20	55% 100% 100%
40	8% H ₂ O ₂ + 1% Acetic Acid Ph 2.6	10 20 30	10/20 8/20 0/20	50% 60% 100%
45	8% H ₂ O ₂ + 0.5% Acetic Acid pH 2.7	10 20 30	11/20 9/20 0/20	45% 55% 100%
50	2% Acetic Acid pH 2.7	30	20/20	Zero

Tests were done at ambient conditions comparing the rapid sporicidal activity of hydrogen peroxide in combination with the carboxylic acids acetic, malonic, succinic, glutaric and citric acids. Compositions from the data are reported in Table III.

TABLE III

Sterilization of *C.sporogenes*-labeled porcelain cylinders by formulations of H₂O₂ plus acetic, malonic, or succinic acid.

Formulation	Exposure Time Min. x 20°C	Percentage of Twenty <i>C.sporogenes</i> -labeled Cylinders Sterilized
8% H ₂ O ₂ plus 1% acetic acid pH 2.5	10 20 30	50% 80 100
8% H ₂ O ₂ plus 1% malonic acid pH 1.8	10 20 30	Zero % 40 85
8% H ₂ O ₂ plus 0.5% malonic acid pH 1.9	10 20 30	5% 25 100
8% H ₂ O ₂ plus 1% succinic acid pH 2.4	10 20 30	Zero % 15 95

Further tests combined H₂O₂ with glutaric acid and citric acid. The results are shown in Table IV and V.

TABLE IV

Sterilization of *C.sporogenes*-labeled porcelain cylinders by formulations of H₂O₂ plus acetic, glutaric, and citric acid.

5	Formulation	pH Value	Exposure Time Min. x 20°C	Percentage of Twenty <i>C.sporogenes</i> -labeled cylinders Sterilized
10	8% H ₂ O ₂ plus 0.2M Acetic Acid	2.4	20	100%
			30	100
15		4.3	20	100%
			30	100
20	8% H ₂ O ₂ plus 0.2 M Glutaric Acid	2.2	20	100
			30	100
		5.0	20	Zero
			30	Zero
25		1.9	20	Zero
			30	Zero
30	8% H ₂ O ₂ plus 0.2M Citric Acid	6.6	20	Zero
			30	Zero

TABLE V

Surviving Colonies of Wet Spores of *B.subtilis* After Exposure to Formulations of H₂O₂ Plus Acetic Acid, Glutaric Acid, or Citric Acid.

Formulation	pH Value	Exposure	Surviving Colonies of					
		Time	<i>B.subtilis</i> at Dilution Factors					
		Min.	x 20°	5x10 ¹	5x10 ²	5x10 ³	5x10 ⁴	5x10 ⁵
8% H ₂ O ₂ plus 0.2 M Acetic Acid	2.8	15		19	1	Zero	Zero	
		30	Zero	Zero	Zero			
		60	Zero	Zero	Zero			
	4.4	15		236	121	26	1	
		30	Zero	Zero	Zero			
		60	Zero	Zero	Zero			
	8% H ₂ O ₂ plus 0.2M Glutaric	2.6	15	CONF	CONF	CONF	265	70
			30		25	13		
			60	Zero	Zero	Zero		
5.0		15	CONF	CONF	CONF	TNTC	101	
		30		TNTC	206	39		
		60	Zero	Zero	Zero			
8% H ₂ O ₂ plus 0.2M Citric	2.0	15	CONF	CONF	CONF	TNTC	198	
		30	CONF	CONF	CONF	TNTC		
		60	CONF	CONF	TNTC			
	6.4	15	CONF	CONF	CONF	CONF	283	
		30	CONF	CONF	CONF	CONF		
		60	CONF	CONF	CONF			

CONF = Confluent = in excess of 1000 colonies all touching together (confluent).

TNTC = Too Numerous to Count = 300-1000 colonies/plate.

The following example in Table VI compares the rate of kill of *B.subtilis* spores by formulations of H₂O₂ plus acetic, malonic, or succinic acid. The test method was wet spores of *B.subtilis* in suspension (not on carriers). This is a quantitative test that allows comparison of formulations with more precision than a qualitative (sterile or not

TABLE VI

Formula Number	Formula Description	pH	D-values in Min.*
5			
1.	8% H ₂ O ₂ + 0.2M (1.2%) acetic acid,	pH 2.7	7.5
2.	8% H ₂ O ₂ + 0.2M (1.2%) acetic acid,	pH 4.2	8.5
3.	8% H ₂ O ₂ + 0.2M (2.1%) malonic acid,	pH 1.8	8.5
10 4.	8% H ₂ O ₂ + 0.2M (2.1%) malonic acid,	pH 3.0	7.8
5.	8% H ₂ O ₂ + 0.2M (2.4%) succinic acid,	pH 2.4	6.0
6.	8% H ₂ O ₂ + 0.2M (2.4%) succinic acid,	pH 4.2	9.0
15	*The D-values were calculated as the time to kill four log ₁₀ of <i>B.subtilis</i> divided by four.		

The general conclusion is that combinations of 8% H₂O₂ plus acetic, malonic, or succinic acid surprisingly have about the same rate of kill of *B.subtilis* spores (wet) in suspension. The more acid pH values of about 2-3 were consistently killing faster than the less acid pH values of above 4.

The tests shown in Tables VII and VIII measure the relationship between increasing concentrations of acetic or succinic acid plus 8% H₂O₂ and the rate of kill of wet spores of *B.subtilis*. The test using suspensions of wet spores of *B.subtilis*, and measuring surviving spores as a function of exposure time to various formulations is a quantitative test that is better able to measure small differences between formulations than the AOAC Sporicidal Test. All tests were at 20±1°C.

Acetic Acid:

The formulations tested with acetic acid, and D-value results were as follows:

TABLE VII

Formula Number	Formula Description	pH Value	D-value
5			
1.	8% H ₂ O ₂ 1.0M (6%) Acetic Acid 0.5% BioTerge AS-40	4.2	less than 3 Min.
10			
2.	8% H ₂ O ₂ 0.5M (3%) Acetic Acid 0.5% BioTerge AS-40	4.2	3.5 Min.
15			
3.	8% H ₂ O ₂ 0.25M (1.5%) Acetic Acid 0.5% BioTerge AS-40	4.3	3.75 Min.
20			
4.	8% H ₂ O ₂ 0.125M (0.75%) Acetic Acid 0.5% BioTerge AS-40	4.3	4.0 Min.

25 BioTerge is a trademark of Stepan Company and is a sodium olefin sulfonate

The formulations tested with succinic acid, and D-value results were as follows in Table VIII:

TABLE VIII

Formula Number	Formula Description	pH Value	D-value
35			
1.	8% H ₂ O ₂ 1.0M (11.8%) Succinic Acid 0.5% BioTerge AS-40	4.3	less than 3 Min.
40			
2.	8% H ₂ O ₂ 0.5M (5.9%) Succinic Acid 0.5% BioTerge AS-40	4.2	3.5 Min.
45			
3.	8% H ₂ O ₂ 0.25M (2.95%) Succinic Acid 0.5% BioTerge AS-40	4.2	3.5 Min.
50			
4.	8% H ₂ O ₂ 0.125M (1.47%) Succinic Acid 0.5% BioTerge AS-40	4.2	3.0 Min.

As demonstrated in Tables VII and VIII at equal molarities, and equal pH values of about 4.2, there is very little difference between acetic acid and succinic acid to enhance spore kill in combination with 8% H₂O₂ and BioTerge AS-40.

5 As one covers a range from a high of 1.0M to a low of 0.125M, an eight-fold difference, the rate of spore kill changes very little from the slowest rate of about 4.0 min. to the fastest rate of about 3.0 min. The change is consistent with 1.0M acetic or succinic acid always showing a
10 faster spore kill than lesser concentrations, but it is a very small change.

8% H₂O₂ (2.35M) plus 1% (0.17M) acetic acid, or plus 0.2M acetic acid at pH 2.5 (all of the acetic acid in the acid form), or pH - 4.3 (about half of the acetic acid in the
15 acid form, and half as sodium acetate) killed spores of *C.sporogenes* and *B.subtilis* within 30 min. at 20±1°C. More *B.subtilis* spores were killed within 15 min. at the lower pH (about 2.5) than at the higher pH (about 4.3), but complete kill happened within 30 min. for both pH values.

20 8% H₂O₂ plus 0.5% malonic acid or 0.5% succinic acid
 COOH-CH₂-COOH COOH-CH₂-CH₂-COOH
 Malonic Acid Succinic Acid

all at pH 1.8 - 2.4 sterilized porcelain cylinders labeled
25 with *C.sporogenes* within 30 min. at 20±1°C.

8% H₂O₂ plus 0.2M glutaric acid (COOH-CH₂-CH₂-CH₂-COOH) was less active than succinic acid, and more active than citric acid. 8% H₂O₂ plus 0.2M glutaric acid at pH 2.2 - 2.6
30 killed wet *B.subtilis* spores within 60 min. at 20±1°C, but not within 30 min., and sterilized *C.sporogenes* dried onto porcelain cylinders within 30 min. at 20±1°C, pH 2.2, but not at pH 5.0 (where half of the glutaric acid exists as the acid and half exists as the sodium salt).

8% H₂O₂ plus 0.2M citric acid, a tricarboxylic acid,
35 COOH-CH₂-COHCOOH-CH₂-COOH, did not kill wet *B.subtilis* spores at all within 60 min. at 20±1°C either at pH 2.0 or 6.4. Similarly, 8% H₂O₂ plus 0.2M citric acid at pH 1.9 or 6.6

could not sterilize any *C.sporogenes*-labeled cylinders at 20±1°C.

The above tests lead to the following observations:

The initial testing was performed because of the similar structural chemistry between peracetic acid ($\text{CH}_3\text{-COOOH}$) and acetic acid ($\text{CH}_3\text{-COOH}$). The following formulations were tested: (1) 6% H_2O_2 , pH 4.7, (2) 6% H_2O_2 + 0.5% acetic acid, pH 2.7, (3) 6% H_2O_2 + 0.5% sodium acetate, pH 6.7, (4) 0.5% acetic acid, pH 2.8, and (5) 0.5% sodium acetate, pH 7.7. Unglazed porcelain cylinders labeled with approx. 10^6 dry spores of *Clostridium sporogenes* according to the methods of the AOAC Sporicidal Test 966.04 were exposed to these above five formulations for 30 min. at 20±1°C. Of a total of 20 spore-labeled cylinders, 18 were sterilized when exposed to 6% H_2O_2 + 0.5% acetic acid (Formula #2 above). None (zero) of 20 were sterilized when similarly exposed to the other formulas.

Two cylinders from a total of 20 tested were not sterilized by formula #2 above. Next tested were increased concentrations of H_2O_2 and acetic acid for ability to sterilize dry cylinders labeled with *C.sporogenes*. 8% H_2O_2 plus 2% acetic acid sterilized within 20 min. at 20±1°C, 8% H_2O_2 plus 1% acetic acid, and also 8% H_2O_2 plus 0.5% acetic acid sterilized within 30 min. at 20±1°C.

The above tests were then expanded to determine if other weak di- or poly carboxylic acids could act synergistically with H_2O_2 for rapid sporicidal activity. The dicarboxylic acids, malonic ($\text{COOH-CH}_2\text{-COOH}$), and succinic ($\text{COOH-CH}_2\text{-CH}_2\text{-COOH}$) were tested. 8% H_2O_2 plus 1% acetic acid, pH 2.5, sterilized within 30 min. at 20±1°C. 8% H_2O_2 plus 1% malonic acid, pH 1.8, sterilized 17 cylinders out of 20 within 30 min. at 20±1°C, and 8% H_2O_2 plus 0.5% succinic acid pH 2.4, sterilized 19 cylinders out of 20 within 30 min. at 20±1°C. The test was *C.sporogenes* on porcelain cylinders.

H_2O_2 plus the weak carboxylic acids produce a formula with acidic pH values of 2-3. However, the carboxylic acids can act as buffers by taking them up to a pH value where half

of the molecules are in the acid form ($\text{CH}_3\text{-COOH}$) and half of the molecules are the conjugate base ($\text{CH}_3\text{-COO}^-$). These higher pH values (pH 4-5) might be more compatible with materials than the lower pH values. 8% H_2O_2 plus 0.2 M citric acid (a tricarboxylic acid) did not sterilize *C.sporogenes*-labeled cylinders at pH 1.9 or 6.6. 8% H_2O_2 plus 0.2 M glutaric acid ($\text{COOH-CH}_2\text{-CH}_2\text{-CH}_2\text{-COOH}$) did sterilize within 30 min. at $20\pm 1^\circ\text{C}$ at pH 2.2, but not at pH 5.0. 8% H_2O_2 plus 0.2 M succinic acid ($\text{COOH-CH}_2\text{-CH}_2\text{-COOH}$) sterilized within 30 min. at $20\pm 1^\circ\text{C}$ at both pH 1.8 and 3.0. 8% H_2O_2 plus 0.2 M acetic acid sterilized within 30 min. at $20\pm 1^\circ\text{C}$ at both pH 2.7 and 4.2. These studies were all against *C.sporogenes*-labeled cylinders.

Studies have been performed using these formulations against *Bacillus subtilis* spores both in the form of dry spores carried on porcelain cylinders, or wet spores in suspension. The results against *B.subtilis* are consistent with results against *C.sporogenes* as follows: 8% H_2O_2 plus 0.2 M citric acid at pH 2.0 or 6.4 did not kill wet spores of *B.subtilis* within 60 min. at $20\pm 1^\circ\text{C}$. 8% H_2O_2 plus 0.2 M acetic acid (pH 2.7 or pH 4.2) killed wet or dry *B.subtilis* within 30 min. at $20\pm 1^\circ\text{C}$. 8% H_2O_2 + 0.2 M malonic killed wet spores of *B.subtilis* faster at pH 1.8 than at pH 3.0. 8% H_2O_2 + 0.2 M succinic acid at pH 2.4 killed wet or dry spores of *B.subtilis*, but was less effective at pH 4.2. 8% H_2O_2 plus 0.2 M glutaric acid was more effective against wet spores of *B.subtilis* at pH 2.6 than at pH 5.0. The specific test results are described below.

This example compares the rate of kill of *B.subtilis* spores by formulations of H_2O_2 plus acetic, malonic, or succinic acid. The test method utilized wet spores of *B.subtilis* in suspension (not on carriers). This is a quantitative test that allows comparison of formulations with more precision than a qualitative (sterile or not sterile) test such as the AOAC Sporocidal Test. The starting number of cells of *B.subtilis* was very high at about 3.1×10^8 cells.

It required about 60 min. of exposure time at 20±1°C to kill all of these cells.

The results were as follows:

5

TABLE IX

Formula Number	Formula Description	pH	D-values in Min.
10	1. 8% H ₂ O ₂ + 0.2M (1.2%) acetic acid,	pH 2.7	7.5
	2. 8% H ₂ O ₂ + 0.2M (1.2%) acetic acid,	pH 4.2	8.5
	3. 8% H ₂ O ₂ + 0.2M (2.1%) malonic acid,	pH 1.8	8.5
	4. 8% H ₂ O ₂ + 0.2M (2.1%) malonic acid,	pH 3.0	7.8
	5. 8% H ₂ O ₂ + 0.2M (2.4%) succinic acid,	pH 2.4	6.0
15	6. 8% H ₂ O ₂ + 0.2M (2.4%) succinic acid,	pH 4.2	9.0

The D-values were calculated as the time to kill four log₁₀ of *B.subtilis* divided by four.

- 20 The general conclusion is that combinations of 8% H₂O₂ plus acetic, malonic, or succinic acid have about the same rate of kill of *B.subtilis* spores (wet) in suspension. The more acid pH values of about 2-3 were consistently killing faster than the less acid pH values of about 4.
- 25 Next, three formulations of 8% H₂O₂ (a pH 2.00 succinic acid formulation, a pH 4.35 succinic acid formulation, and a pH 4.23 acetic acid formulation) were placed into plastic trays with loose-fitting plastic lids. Various combinations of stainless steel instruments, endoscope parts, and
- 30 respiratory care equipment were soaked in the formulations for fourteen days at ambient temperature (22±2°C). Two marketed disinfectants (2% alkaline glutaraldehyde, and 0.25% quaternary ammonium compounds in 15% isopropanol) were also used in the study for comparison.
- 35 After fourteen days of continuous soaking in the H₂O₂ formulations, quality Sklarlite® stainless steel instruments appeared unchanged. Less expensive, poorly-plated instruments became mildly tarnished by the three H₂O₂ formulations. By comparison, the quality Sklarlite®
- 40 instruments had become slightly rusted by 2% alkaline glutaraldehyde and extremely rusted by the alcohol

disinfectant. With one exception, the endoscope parts and respiratory care equipment appeared unchanged by any of the H₂O₂ + carboxylic acid formulations. The details of the test are reported below.

- 5 The study of this example was limited to visual observations of materials compatibility with the formulations as previously described. In particular, the formulations used were:

10 Formulation #1

8% H₂O₂

0.5 M Acetic Acid

0.25% Bio-Terge AS-40 detergent

0.25 M NaOH

- 15 Prepared with USP purified deionized H₂O
pH 4.23

Formulation #2

8% H₂O₂

- 20 0.5 M Succinic Acid

0.25% Bio-Terge AS-40 detergent

0.5 M NaOH

Prepared with USP purified deionized H₂O
pH 4.35

25

Formulation #3

8% H₂O₂

0.5 M Succinic Acid

0.25% Bio-Terge AS-40 detergent

- 30 Prepared with USP purified deionized H₂O
pH 2.00

The materials were soaked in the above test formulas or in:

- 35 0.25% quaternary ammonium chloride in 15% isopropanol, or
2% alkaline glutaraldehyde.

The items soaked were:

Eight Cambro plastic trays with loose-fitting plastic lids;

- 5 Five Sklarlite® stainless steel Halsted Mosq. STR 5"
Hemostats. Sklar Hospital Catalog #23-2105. New;

Three pair of inexpensive scissors, poorly plated, but
otherwise in good condition with no tarnish;

10

One set of respiratory care equipment:

- a "Y" plastic connector
a face mask
15 an endotracheal tube
a section of a blue latex breathing bag

Two sets of endoscope parts, the first being:

- insertion tube, bending rubber, biopsy channel, pliable ≈
20 1/3" id connector, hard ≈ 1/2" id connector, and hard ≈ 1/2"
diam. cap.

The second set was an insertion tube, bending rubber, biopsy
channel, hard ≈ 1/2" id connector, hard ≈ 1/2" diam. cap, and
25 hard ≈ 1/2" diam. cap with stainless steel opening.

All parts were new or in good condition at the start.

Two hundred ml of disinfectant and various instruments,
30 parts, and equipment were placed into eight plastic trays.
The trays were covered and left at ambient temperature (22±
2°C) for fourteen days. Observations were made at various
intervals throughout the fourteen day time. The results are
reported in Table X, below.

35

Results:

TABLE X

Observations of Materials Compatibility
Exposure Time to Disinfectant

40

<u>Disinfectant</u>	<u>Instrument</u>	<u>Day 2</u>	<u>Day 3</u>	<u>Day 6</u>	<u>Day 9</u>	<u>Day 14</u>
2% Alkaline glutaraldehyde	Hemostats	N.C.	N.C.	N.C.	slight rust in hinge	slight rust in hinge

0.25% quaternary ammonium chloride in 15% isopropanol	Hemostats	N.C.	2-3 mm rust spot in hinge	major rust in hinge	major rust in hinge	major rust in hinge
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Formulation #1

8% H ₂ O ₂ 0.5 M Acetic Acid	Hemostats	N.C.	N.C.	N.C.	N.C.	N.C.
0.25% Bio- Terge AS-40 0.25M NaOH pH = 4.23	Scissors	N.C.	N.C.	Mild tarnish	Tarnish on handle & hinges	Tarnish on handle & hinges

Formulation #3

8% H ₂ O ₂ 0.5 M Succinic Acid	Hemostats	N.C.	N.C.	N.C.	N.C.	N.C.
0.25% Bio- Terge AS-40 pH = 2.00	Scissors	N.C.	N.C.	Mild tarnish	Tarnish on handle & hinges	Tarnish on handle & hinges

Formulation #2

8% H ₂ O ₂ 0.5 M Succinic Acid	Hemostats	N.C.	N.C.	N.C.	N.C.	N.C.
0.25% Bio- Terge AS-40 0.5 M NaOH pH = 4.35	Scissors	N.C.	slight tarnish in hinge	Mild tarnish	Tarnish on handle & hinges	Tarnish on handle & hinges

Formulation #2

8% H ₂ O ₂ 0.5 M Succinic Acid	"y"	N.C.	N.C.	N.C.	N.C.	N.C.
0.25% Bio- Terge AS-40 0.5 M NaOH pH = 4.35	Connector Face Mask Endo- tracheal Tube Breathing Tube	N.C. N.C. N.C. N.C. N.C.	N.C. N.C. N.C. N.C. N.C.	N.C. N.C. N.C. N.C. N.C.	N.C. N.C. N.C. N.C. N.C.	N.C. N.C. N.C. N.C. N.C.

<u>Formulation #3</u>		Insertion	N.C.	N.C.	N.C.	N.C.	N.C.
8% H ₂ O ₂	Tube						
0.5 M Succinic Acid	Biopsy Channel	N.C.	N.C.	N.C.	N.C.	N.C.	
0.25% Bio-Terge AS-40	Bending Rubber	N.C.	N.C.	N.C.	N.C.	N.C.	
pH = 2.00	Pliable Connector	N.C.	N.C.	N.C.	N.C.	N.C.	Broke into small pieces
	Hard Connector	N.C.	N.C.	N.C.	N.C.	N.C.	N.C.
	Hard Cap	N.C.	N.C.	N.C.	N.C.	N.C.	N.C.
<u>Formulation #2</u>							
8% H ₂ O ₂	Insertion	N.C.	N.C.	N.C.	N.C.	N.C.	
0.5 M Succinic Acid	Tube						
0.25% Bio-Terge AS-40	Biopsy Channel	N.C.	N.C.	N.C.	N.C.	N.C.	
0.5 M NaOH	Bending Rubber	N.C.	N.C.	N.C.	N.C.	N.C.	
pH = 4.35	Hard Cap	N.C.	N.C.	N.C.	N.C.	N.C.	
	Hard Connector	N.C.	N.C.	N.C.	N.C.	N.C.	
	Hard Cap with stainless steel opening	N.C.	N.C.	N.C.	N.C.	N.C.	

N.C. = No Change

5 As seen from data in Table X, formulations 1, 2 and 3 did not cause any apparent changes to the quality Sklarlite® instruments. The formulations did cause some tarnishing of the poorly-plated instruments. The pH 2.00 8% H₂O₂, succinic acid formulation caused more tarnishing than the other two
10 formulations.

In comparison, 2% alkaline gluteraldehyde caused minor rusting of the quality Sklarlite® hemostats, and 0.25% quaternary ammonium chloride in 15% isopropanol caused major rusting of the Sklarlite® hemostats.

15 The pH 2.00, 8% H₂O₂, succinic acid formulation did cause major disintegration of one piece, the pliable endoscope connector, which fell apart when squeezed slightly. It was not known whether this aberation was caused by the nature of the elastomer of this single part or not. However,

no other parts during testing showed any damage by succinic acid compositions.

Formulations 2 and 3 did not cause any apparent change to the other endoscope parts. Formulation 2 did not cause
5 any apparent change to the respiratory care equipment.

While not wishing to be bound by a theory of why the invention works, the data in the above examples demonstrates an apparent reaction and a synergistic relationship between hydrogen peroxide specifically and certain of the described
10 carboxylic acids. It probably extends to peroxides in general that release hydroxyl free radicals that together cause rapid kill of bacterial spores and all other microbes at ambient (approximately 18°C-24°C) temperatures. There is no need for heating, and moreover the kill is generally
15 accomplished within 30 min. It also suggests that a reaction product may be formed in situ which could be isolated and itself used as the quick acting sterilant, and thus the invention contemplates such an embodiment as being within its scope.

20 It therefore can be seen that the invention accomplishes all of its stated objectives.

What is claimed is:

1. A low odor, aqueous quick acting room temperature
disinfecting and/or sterilization solution having a pH within
5 the range of from about 2.0 to about 6.0, comprising: from
about 1% to about 30% by weight of a peroxide; and from about
1% to about 30% by weight of water soluble organic acid or
salt form thereof, selected from the group consisting of C₃
or higher mono carboxylic acids and di- or poly carboxylic
10 acids of up to C₁₂ chain length, or mixtures thereof.

2. The aqueous disinfecting and/or sterilizing solutions of
claim 1 wherein the water soluble organic acid or salt form
thereof is selected from the group consisting of malonic
15 acid, and succinic acid, or mixtures thereof.

3. An aqueous disinfecting and/or sterilizing solution of
claim 2 wherein the peroxide has a concentration of from
about 1.0% by weight to about 12% by weight.
20

4. An aqueous disinfecting and/or sterilization solution of
claim 1 wherein the peroxide capable of releasing hydroxyl
free radicals is selected from the group consisting of
hydrogen peroxide, alkyl peroxides, aryl peroxides, ozonides,
25 and alkylidene peroxides.

5. A low odor aqueous quick acting, relatively non-toxic
room temperature disinfecting and/or sterilization solution
having a pH within the range of from about 2.0 to about 6.0,
30 comprising: from about 1.0% to about 30% by weight of a
peroxide capable of releasing hydroxyl free radicals; from
about 1.0% to about 30% by weight of a water soluble organic
acid or a salt form thereof, selected from the group
consisting of C₃ or higher mono and dicarboxylic acids of up
35 to C₁₂ chain length; and from about 0.1% to about 1.0% by
weight of a peroxide and organic acid compatible anionic or
nonionic surfactant.

6. The aqueous disinfecting and/or sterilizing solution of claim 5 wherein the water soluble organic acid or salt form thereof is selected from the group consisting of malonic acid and succinic acid or mixtures thereof.

7. A process of quick action room temperature disinfecting of medical instruments without damaging the instruments, comprising: contacting at room temperature the instruments for a sterilizing effective amount of time with an odor-free aqueous disinfecting solution having a pH within the range of from about 2.0 to about 6.0 which consists essentially of from about 1.0% by weight to about 30.0% by weight of hydrogen peroxide and from about 1.0% by weight to 30.0% by weight of a low odor water soluble organic acid or a salt form thereof selected from the group consisting of malonic and succinic acids or mixtures thereof.

8. The process of claim 7 wherein the organic acid is succinic acid.

9. The process of claim 7 wherein the aqueous disinfecting solution contains a peroxide and organic acid compatible surfactant.

INTERNATIONAL SEARCH REPORT

National Application No
PCT/US 97/02183

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 A01N59/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 A01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DE 26 54 164 A (SCHÜLKE & MAYR) 22 December 1977 *see the whole document*	1-9
P, X	DATABASE WPI Derwent Publications Ltd., London, GB; AN 96-196557 XP002033260 see abstract & JP 08 067 667 A	1-9
X	CHEMICAL ABSTRACTS, vol. 114, no. 2, 1991 Columbus, Ohio, US; abstract no. 12263, XP002033259 see abstract & SU 1 572 478 A	1-9

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☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 97/02183

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DE 43 31 942 A (KARL LÖFFLER GMBH) 23 March 1995 see column 1, line 58 - column 4, line 10 see examples 1,2,6 ---	1,4,5
X	WO 93 05016 A (PREPARATION OF PEROXYACIDS) 18 March 1993 *see the whole document* ---	1,4,5
X	WO 91 08981 A (AQUACLEAR INTERNATIONAL LIMITED) 27 June 1991 see page 5, line 15 - page 9, line 20 see examples 1-5 ---	1,4,5
X	EP 0 423 922 A (ECOLAB INC.) 24 April 1991 * see the whole document* -----	1,4,5

INTERNATIONAL SEARCH REPORT

Information on patent family members

National Application No

PCT/US 97/02183

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